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(54) Title: EBV CTL EPITOPES (57) Abstract <p>The present invention provides cytotoxic Epstein-Barr virus T-cell epitopes. These epitopes are QVKWRMTTL, VFSDGRVAC, VPAPAGPIV, TYSAGIVQI, LLDFVRFMGV, QNGALAINTF, VSSDGRVAC, VSSEGRVAC, VSSDGRVPC, VSSDGLVAC, VSSDGQVAC, VSSDGRVVC, VPAPPVGPIV, VEITPYEPTG, VEITPYEPTW, VELTPYKPTW, RRIYDLIKL, RKIYDLIEL and PYLFWLAGI. The present invention further provides vaccines including one or more of these epitopes, optionally with additional epitopes.</p>		

EBV CTL Epitopes**FIELD OF THE INVENTION**

The present invention relates to cytotoxic T-cell (CTL) epitopes
5 within Epstein-Barr virus. The present invention also relates to subunit
vaccines including CTL epitopes.

BACKGROUND OF THE INVENTION

Epstein-Barr virus (EBV) is a gamma herpesvirus which establishes a
10 latent lifelong infection in the host following acute infection (14,15). While
primary infection generally occurs in childhood without significant
morbidity, adolescents and young adults may present with the symptoms of
acute infectious mononucleosis (IM). The main feature of IM is a self-
limiting lymphoproliferation involving both T and B cells accompanied by
15 clinical symptoms such as fever and lymphadenopathy (52,53).
Occasionally, the clinical symptoms persist and recur for extended periods
after the initial infection. Episodic IM such as this has been described as
chronic active EBV infection or, in some cases, severe chronic active EBV
infection (35). EBV DNA has been detected in both serum and peripheral
20 blood lymphocytes (PBL) during acute IM with the levels of detectable DNA
gradually decreasing as the illness abates (21,22,58).

Evidence for latent EBV infection includes the observation that
spontaneous lymphoblastoid cell lines (LCLs), expressing latent proteins,
can be regularly established from healthy immune individuals after
25 explantation of either lymph node tissue (34) or fractionated B lymphocytes
(59). Although latent EBV infection is usually asymptomatic, sequential
studies have established that recrudescence of viral replication in the oral
cavity may result in release of infectious virus (59). The exact site of
persistence of the virus is uncertain, but the available evidence suggests that
30 small lymphocytes in the circulation harbour the virus in a nonproductive
episomal state (24). Accordingly, in asymptomatic donors, EBV DNA is
detectable by sensitive PCR analysis in PBL expressing the B-cell marker
CD19 (29,55).

EBV is also involved in post transplant lymphoproliferative disease.
35 which involves a polyclonal expansion of EBV infected B-cells which is a

life threatening lymphoma especially in transplantation patients. EBV is also involved in nasopharyngeal carcinoma and Hodgkinson's disease.

Two types of EBV (types A and B or types 1 and 2) are distinguishable primarily on the basis of variation in the DNA and protein sequences of the latent EBV-associated nuclear antigens (referred to here as EBNA-2A, -3A, -4A and -6A from the type A virus and EBNA-2B, -3B, -4B and -6B from the type B virus) (9,46,48). Sequencing of the prototypical isolates of type A and type B EBV (B95-8 and Ag-876 respectively) in these regions revealed 53% amino acid homology between EBNA-2A and EBNA-2B (9) and 72-84% homology between EBNA-3A and -3B, EBNA-4A and -4B, and EBNA-6A and -6B (46). Strain variation due to other DNA alterations or deletions as well as these A/B type differences have been defined at the protein (12) and the DNA level (16,25,26) and recombination between multiple infecting strains was found to occur frequently in oral hairy leukoplakia lesions (56). These variations offer an alternative means of categorising EBV isolates but the primary distinction of type A and type B is still useful. Type A EBV is more readily isolated from healthy donors; type B EBV infections or dual infections with both type A and type B have proven easier to detect in immunosuppressed or HIV infected individuals (5,47,51). A higher incidence of type B infection in some studies led to the suggestion that type B or dual infections are, in fact, relatively common and that resident type B virus levels increase during immunosuppression (3,20,51).

It appears that latent EBV infection is primarily controlled by HLA class I- restricted memory cytotoxic T cell (CTL) responses (reviewed in (18)). These CTL responses can be reactivated in vitro by stimulating lymphocytes from seropositive individuals with autologous lymphoblastoid cell lines (LCLs) which express and present MHC class I and class II restricted epitopes at the cell surface. Several of these epitopes have been identified using target cells infected by recombinant vaccinia constructs (17,19,32,33). Epitopes specific for type A EBV as well as cross-reactive epitopes encoded by both types A and B EBV have been defined (18) but no epitopes specific for type B EBV have been reported thus far. In addition co-pending International Patent Application No. WO 95/001400, the disclosure of which is incorporated herein by cross reference discloses a number of EBV CTL epitopes. In the present study, the response of a donor exposed to

both type A and type B EBV was investigated and an epitope specific for type B EBV as well as a new cross-reactive epitope were identified.

SUMMARY OF THE INVENTION

5 In a first aspect the present invention consists in a cytotoxic Epstein-Barr virus T-cell epitope, the epitope being selected from the group consisting of QVKWRMTTL, VFSDGRVAC, VPAPAGPIV, TYSAGIVQI, LLDFVRFMGV, QNGALAINTF, VSSDGRVAC, VSSEGRVAC, VSSDGRVPC, VSSDGLVAC, VSSDGQVAC, VSSDGRVVC, VPAPPVGPIV, VEITPYEPTG,
10 VEITPYEPTW, VELTPYKPTW, RRIYDLIKL, RKIYDLIEL and PYLFWLAGI.

In a second aspect the present invention consists in a subunit vaccine, the vaccine including at least one T-cell epitope selected from the group consisting of QVKWRMTTL, VFSDGRVAC, VPAPAGPIV, TYSAGIVQI, LLDFVRFMGV, QNGALAINTF, VSSDGRVAC, VSSEGRVAC, VSSDGRVPC,
15 VSSDGLVAC, VSSDGQVAC, VSSDGRVVC, VPAPPVGPIV, VEITPYEPTG, VEITPYEPTW, VELTPYKPTW, RRIYDLIKL, RKIYDLIEL and PYLFWLAGI.

In a preferred embodiment of this aspect of the present invention the subunit vaccine includes at least one further epitope selected from the group consisting of TSLYNLRRGTALA, DTPLIPLTIF, TVFYNIPPMPL,
20 VEITPYKPTW, VSFIEFVGW, FRKAQIQGL, FLRGRAYGL, QAKWRLQTL, SVRDRRLARL, YPLHEQHGM, HLAAQGMAY, RPPIFIRRL, RLRAEAGVK, IVTDFSVIK, AVFDRKSDAK, NPTQAPVIQLVHAVY, LPGPQVTAVLLHEES, DEPASTEPVHDQLL, RYSIFFDY, AVLLHEESM, RRARSLAERY, EENLLDFVRF, KEHVIQNAF, RRIYDLIEL, QPRAPIRPI, EGGVGWRHW,
25 CLGGLTMTV, RRRWRRLTV, RAKFKQLL and RKCCRAKFKQLQHYP.

In a further preferred embodiment of this aspect of the present invention the subunit vaccine includes the cytotoxic T-cell epitopes LLDFVRFMGV, QVKWRMTTL and FLRGRAYGL. An analysis of allele frequency in the HLA listings in the Queensland Institute of Medical
30 Research data base shows that a vaccine including these epitopes would provide protection for 63.7% of the caucasian population.

In a yet further preferred embodiment of this aspect of the present invention the subunit vaccine includes the cytotoxic T-cell epitopes LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL and EENLLDFVRF. An
35 analysis of allele frequency in the HLA listings in the Queensland Institute

of Medical Research data base shows that a vaccine including these epitopes would provide protection for 71.1% of the caucasian population.

In another preferred embodiment of this aspect of the present invention the subunit vaccine includes the cytotoxic T-cell epitopes
5 LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL and QPRAPIRPI. An analysis of allele frequency in the HLA listings in the Queensland Institute of Medical Research data base shows that a vaccine including these epitopes would provide protection for 74.1% of the caucasian population.

In a still further preferred embodiment of this aspect of the present
10 invention the subunit vaccine includes the cytotoxic T-cell epitopes LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL, EENLLDFVRF and QPRAPIRPI. An analysis of allele frequency in the HLA listings in the Queensland Institute of Medical Research data base shows that a vaccine including these epitopes would provide protection for 81.5% of the
15 caucasian population. Given the fact that about 50% of all individuals that are not covered by vaccination will become EBV positive without any symptoms, the combination of epitopes listed above will result in a vaccine with more than 90% efficacy. This is of high commercial value.

In a further preferred form of the present invention the vaccine
20 comprises a water-in-oil formulation. It is further preferred that the vaccine includes at least one antigen to which the individual will mount an anamnestic response in addition to the at least one cytotoxic T-cell epitope.

The at least one antigen is preferably selected from the group consisting of tetanus toxoid, diphtheria toxoid, Bordetella pertussis antigens,
25 poliovirus antigens, purified protein derivative (PPD), gp350 protein (Thorley-Lawson, D.A. and Poodry, C.A. (1982). Identification and isolation of the main component (gp350-gp220) of Epstein-Barr virus responsible for generating neutralizing antibodies in vivo. J. Virol. 43. 730-736), helper epitopes and combinations thereof and is preferably tetanus
30 toxoid.

It is preferred that the water-in-oil formulation is Montanide ISA 720. Additional information regarding this formulation can be found in WO 95/24926, the disclosure of which is incorporated herein by cross reference.

35 The subunit vaccine may also be formulated using ISCOMs. Further information regarding ISCOMs can be found in Australian Patent

Nos. 558258, 590904, 632067, 589915, the disclosures of which are included herein by cross reference.

In a third aspect the present invention consists in a nucleic acid vaccine, the vaccine including a nucleic acid sequence encoding at least one of the cytotoxic T-cell epitopes of the first aspect of the present invention and optionally at least one cytotoxic T-cell epitope selected from the group consisting of TSLYNLRRGTALA, DTPLIPLTIF, TVFYNIPPMPL, VEITPYKPTW, VSFIEVGVW, FRKAQIQGL, FLRGRAYGL, QAKWRLQTL, SVRDLRLARL, YPLHEQHGM, HLAAQGMAY, RPIFIRRL, RLRAEAGVK, IVTDFSVIK, AVFDRKSDAK, NPTQAPVIQLVHAVY, LPGPQVTAVILLHEES, DEPASTEPVHDQLL, RYSIFFDY, AVLLHEESM, RRARSLSAERY, EENLLDFVRF, KEIIVIQNAF, RRIYDLIEL, QPRAPIRPI, EGGVGWRHW, CLGGLLTMV, RRRWRRLTV, RAKFKQLL and RKCCRAKFKQLLQHYR.

Further information regarding nucleic acid vaccines can be found in WO 96/03144 the disclosure of which is incorporated herein by cross reference. As will be appreciated by those skilled in the field the DNA can be delivered using a suitable viral or bacterial vector.

In a preferred embodiment of this aspect of the present invention the nucleic acid sequence encodes LLDFVRFMGV, QVKWRMTTL and FLRGRAYGL, or LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL and EENLLDFVRF, or LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL and QPRAPIRPI, or LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL, EENLLDFVRF and QPRAPIRPI.

The vaccines of the present invention may be used prophylactically or therapeutically.

DETAILED DESCRIPTION OF THE INVENTION

The CTL epitopes may be synthesised using techniques well known to those skilled in this field. For example, the CTL epitopes may be synthesised using solution synthesis or solid phase synthesis as described, for example, in Chapter 9 entitled "Peptide Synthesis" by Atherton and Sheppard which is included in a publication entitled "Synthetic Vaccines" edited by Nicholson and published by Blackwell Scientific Publications. Preferably a solid phase support is utilised which may be polystyrene gel beads wherein the polystyrene may be cross-linked with a small proportion of divinylbenzene (e.g. 1%) which is further swollen by lipophilic solvents

such as dichloromethane or more polar solvents such as dimethylformamide (DMF). The polystyrene may be functionalised with chloromethyl or anionomethyl groups. Alternatively, cross-linked and functionalised polydimethyl-acrylamide gel is used which may be highly solvated and swollen by DMF and other dipolar aprotic solvents. Other supports can be utilised based on polyethylene glycol which is usually grafted or otherwise attached to the surface of inert polystyrene beads. In a preferred form, use may be made of commercial solid supports or resins which are selected from PAL-PEG, PAK-PEG, KA, KR or TGR.

10 In solid state synthesis, use is made of reversible blocking groups which have the dual function of masking unwanted reactivity in the α -amino, carboxy or side chain functional groups and of destroying the dipolar character of amino acids and peptides which render them inactive. Such functional groups can be selected from *t*-butyl esters of the structure RCO-OCMe_3 -CO-NHR which are known as *t*-butoxy carboxyl or ROC derivatives. Use may also be made of the corresponding benzyl esters having the structure $\text{RCO-OCH}_2\text{-C}_6\text{H}_5$ and urethanes having the structure $\text{C}_6\text{H}_5\text{CH}_2\text{O-CO-NHR}$ which are known as the benzyloxycarbonyl or Z-derivatives. Use may also be made of derivatives of fluorenyl methanol and especially the fluorenyl-methoxy carbonyl or Fmoc group. Each of these types of protecting group is capable of independent cleavage in the presence of one other so that frequent use is made, for example, of BOC-benzyl and Fmoc-tertiary butyl protection strategies.

Reference also should be made to a condensing agent to link the amino and carboxy groups of protected amino acids or peptides. This may be done by activating the carboxy group so that it reacts spontaneously with a free primary or secondary amine. Activated esters such as those derived from *p*-nitrophenol and pentafluorophenyl may be used for this purpose. Their reactivity may be increased by addition of catalysts such as 1-hydroxybenzotriazole. Esters of triazine DHBT (as discussed on page 215-216 of the abovementioned Nicholson reference) also may be used. Other acylating species are formed in situ by treatment of the carboxylic acid (i.e. the Na-protected amino acid or peptide) with a condensing reagent and are reacted immediately with the amino component (the carboxy or C-protected amino acid or peptide). Dicyclohexylcarbodiimide, the BOP reagent (referred to on page 216 of the Nicholson reference), O-Benzotriazole-N, N,

N,N'-tetra methyl-uronium hexafluorophosphate (HBTU) and its analogous tetrafluoroborate are frequently used condensing agents.

The attachment of the first amino acid to the solid phase support may be carried out using BOC-amino acids in any suitable manner. In one method BOC amino acids are attached to chloromethyl resin by warming the triethyl ammonium salts with the resin. Fmoc-amino acids may be coupled to the p-alkoxybenzyl alcohol resin in similar manner. Alternatively, use may be made of various linkage agents or "handles" to join the first amino acid to the resin. In this regard, p-hydroxymethyl phenylactic acid linked to aminomethyl polystyrene may be used for this purpose.

Details of the CTL epitopes of the present invention are set in Tables 3 and 4.

As will be readily appreciated by those skilled in the art the cytotoxic T-cell epitopes and vaccines of the present invention can be used to protect against EBV. Further given the possible greater involvement of type B EBV infection in immunocompromised individuals the present invention may have particular application in protection of individuals having decreased immune function, eg transplant patients.

In order that the nature of the present invention may be more clearly understood preferred forms thereof will now be described with reference to the following examples and Figures.

FIGURE LEGENDS

Fig 1. Immunoblot of recombinant vaccinia-EBNA viruses: Lane 1: vacc-TK-, Lane 2: vacc-EBNA-3B; Lane 3: vacc-EBNA-6B. M. Wt. markers are on the left.

Fig. 2. Confirmation of specificity of CTL cultures B5 and A3. EW-Wil (type A autologous LCL) was infected with vacc- EBNA-6A, vacc-EBNA-6B or vacc-TK and used as targets in a CTL assay: EW-Wil and EW-Ag-876 (type B autologous LCL) were used as control targets without vaccinia infection.

Fig. 3a, b: Identification of minimal epitope for culture B5. Targets: EW-Wil LCL + peptide

Fig. 3a: Targets + peptides 7-12 of EBNA 6B 20-mer overlap

Fig. 3b: Targets + 9-mer and 10-mer overlapping peptides of the 20-mer peptide No. 12

Fig. 4a, b: Identification of minimal epitope for culture A3. Targets:
5 EW-PHA blasts + peptide

Fig. 4a: Targets + peptides 19-24 of EBNA-6B 20-mer overlap

Fig. 4b: Targets + dilutions of four of the 10-mer overlapping peptides of the 20-mer peptide No. 19

10 **Fig. 5:** Depletion experiment. Targets: EW-C3 (negative T cell culture); Effectors: EW type B epitope specific cultures pooled from LDA cultures. EW = CTL treated with Dynabeads only; EW-CD-8: CTL depleted of CD4 T cells; EW-CD4: CTL depleted of CD8 T cells

15 **Fig. 6:** Inhibition of type B epitope-positive CTL (culture B-15) by a-Class I but not by a-Class II monoclonal antisera. Peptide 74 = type B reactive peptide; target cells = EW-C3, a noncytolytic T cell culture.

MATERIALS AND METHODS

20 *Lymphocyte donor*

A 17 year old individual (EW) who initially presented with clinical and pathological (EBV IgM positive) symptoms of acute IM was used in this study. She was referred to this laboratory for investigation of her EBV status 7 months after the initial diagnosis as her clinical symptoms persisted. Four
25 bleeds were obtained from this donor over a period of 12 months during which time the symptoms fluctuated but did not completely resolve. The third and fourth bleeds were used for CTL analysis.

Virus Typing and Detection

DNA was extracted from PBL by a small scale adaptation of a
30 published extraction procedure (27). EBNA 2A and 2B primers as previously described (20) were used to distinguish between type A and type B EBV by PCR. Briefly, standard PCR reactions were carried out in a 25 µl volume using 1 µg genomic DNA. The template was denatured for 2 min at 94°C before subjecting to 35 PCR cycles (15 s at 94°C, 30 s at 56°C and 15 s at
35 72°C). PCR amplified products were identified on southern blots using digoxigenin-dUTP (DIG) labelled DNA probes (Boehringer-Mannheim

Australia) generated by PCR using the same primers as above and either the purified M-ABA Bam HI insert from pM-Bam H2 (type A) (36) or the purified Jijoye Bam HI/Pst I insert from pJ-HKA7 (type B) (1) as templates. For the preparation of the probes, DIG-dUTP was present in the PCR reaction with 25 ng of each insert as template.

Vaccinia constructs

Recombinant vaccinia constructs (designated vacc-EBNA-1, vacc-EBNA-2 etc) encoding EBNA-1, -2B, -4A and -6A, the latent membrane proteins LMP1 and LMP2, the lytic antigens BHRF1 and BZLF1, and the control vacc-TK have already been described (19.23.33). Vacc-EBNA-3B and vacc-EBNA-6B (i.e. vaccinia constructs encoding the type B sequences of EBNA 3 and EBNA 6 respectively) were constructed from an EBV genomic library derived from purified viral genomes of the prototype type B strain Ag-876 (50). The viral DNA was digested with HindIII and ligated into the λ -ZAP cloning system (Integrated Sciences).

An 11.07 kbp HindIII fragment encoding the EBNA 3 family open reading frames (ORFs) was subcloned into the vector pUC19. A 2.6 kbp BamHI-CEIII (Klenow-treated) fragment within the region containing the BHRF1 ORF of EBNA-3 (46) was isolated and ligated into a BamHI-HindII digest of pBCB07. This vaccinia vector drives recombinant gene expression by the 7.5K vaccinia virus promoter (2). The EBNA-3B sequence encoded in the transfer plasmid was inserted into vaccinia virus by transfection and homologous recombination into the TK gene of vaccinia as described (2). A 4 kbp Eco47III-HindIII fragment within the EBV HindIII region containing the BHRF2b ORF of EBNA-6B was isolated and ligated into a HindII-HindIII digest of pBCB07. This was subsequently processed as for EBNA-3B.

Integrity of the vacc-EBNA-3B and -6B constructs was confirmed by immunoblot after infection of mouse CV-1 cells. 2×10^5 CV-1 cells infected with control vaccinia (vacc-TK), vacc-3B or vacc-6B were lysed in protein sample buffer and separated on a 7% SDS-polyacrylamide gel in a Mini-protean II Cell System (Bio-Rad). After transferring to nitro-cellulose the blot was incubated with a multivalent human serum (MCR) which contained EBNA-specific antibodies. The reactions were visualised with the ECL detection system (Amersham). Fig. 1 illustrates the production of a 110 kDa protein (EBNA-3B) or a 130 kDa protein (EBNA-6B) corresponding to the

expected products of EBNA-3B and EBNA-6B without the short first ORFs. Sequence analysis also confirmed the integrity of the constructs.

Establishment of cell cultures

- 5 (a) B Cell Cultures: LCLs were established from EW by exogenous transformation of PBL by QIMR-WIL virus (type A; (38)) and Ag-876 (type B). These LCLs were designated EW-WIL and EW AG-876. Spontaneous LCLs (i.e. autologous LCLs infected with the donor's endogenous virus strain and designated EW-spon) were established in the presence of cyclosporin (41).
- 10 LCLs from other donors were transformed by QIMR-WIL, B95-8 or IARC-BL74 virus (30) (type A isolates) or by Ag-876 or QIMR-GOR virus (37) (type B isolates). Other lines used in this study include Burkitt lymphoma cell lines (Chop and Mutu) and the spontaneous line QIMR-WW2-LCL (13.43.44). B cell lines were maintained in RPMI 1640 with 10-20%FCS.
- 15 (b) T Cell Cultures: Unfractionated PBL were separated on Ficoll-Paque (Pharmacia, Uppsala, Sweden) and stimulated by incubating with γ -irradiated (8000 rads) autologous LCLs (EW-Wil or EW-Ag-876; 2×10^6 cells per well in 2 ml growth medium without rIL-2, responder:stimulator ratio 50:1) (28). After 3 days, the cultures were cloned in agarose (Seaplaque,
- 20 FMC Corp. Rockland ME) and the colonies harvested 3-5 days later. Once established, the T cells were maintained in T cell growth medium (RPMI-1640 supplemented with 15% FCS, 30% MLA-144 supernatant (TIB-201: American type Culture Collection, Rockville, MD and rIL-2, 20 units/ml) (45.57). Since the clonal nature of the T cells was not established, they are
- 25 referred to in the text as cultures rather than clones. The cultures were routinely maintained in either 24-well plates or tissue culture flasks and restimulated twice weekly with γ -irradiated (8000 rads) autologous LCLs.
- 30 Bulk T cell cultures were initially stimulated from the fourth bleed as for cloning but subcultured and maintained after 3-4 days as for T cell colonies. Phytohaemagglutinin (PHA) blasts were generated by stimulation of PBL with PHA (CSL, Melbourne, Australia) and subculturing after 4 days in T cell growth medium. Cultures were regularly screened for mycoplasma contamination.

FACS analysis

Three-colour cytofluorographic analysis of T cell cultures was performed by means of direct immunofluorescence using labelled monoclonal antibodies specific for CD3 (tricolour- conjugated), CD4 (fluorescein -conjugated), and CD8 (phycoerythrin-conjugated) and (Becton Dickinson, California). The labelled cells were analysed on a FACScan (Becton Dickinson, lysis II software).

Cytotoxicity assays

T cells were assayed in modified 5-hour ^{51}Cr -release assays (30). Briefly, target cells were labelled for 1 hour with $0.5\text{-}1.0\mu\text{Ci } ^{51}\text{Cr}$, washed and resuspended at 10^5 cells/ml. $50\mu\text{l}$ of target cells were incubated with an equal volume of effectors (Effector:target or E:T ratio generally 10:1) in 96-well V-bottomed microtiter plates, centrifuged, and incubated at 37°C for 5 hours. After centrifuging again, $25\mu\text{l}$ supernatant was harvested and dried onto 96-well solid scintillation microtiter plates before counting the radioactivity in a Topcount Microplate Scintillation Counter (Packard Instrument Company, Meridan, CT).

For determination of antigenic specificity, target cells were infected with vaccinia construct (multiplicity of infection approx. 10:1) and labelled with ^{51}Cr by a modification of the previously described procedure (17). Both infection and labelling occurred simultaneously for 2.5 hours before washing and using the cells in CTL assays. In some instances target cells were pretreated in V-wells with monoclonal antibodies to HLA-class I (W6/32) or class II (L243) for 30 min at room temperature before adding effector cells.

Regression assay

The EBV-specific memory CTL response was assessed as previously described (31).

Limiting dilution cultures

Limiting dilution cultures were established by distributing PBL from the fourth bleed in 24 replicates at 2×10^4 cells per well in 96-well round-bottomed microtitre plates. Doubling dilutions were made to a final concentration of 1.6×10^2 per well. Approximately 1×10^4 γ -irradiated (8000 rads) autologous type B (EW-Ag-876) LCLs were added to each well as both feeder and stimulator cells to a total volume of $100\mu\text{l}$. Cultures were fed on days 4 and 7 with $50\mu\text{l}$ of T cell growth medium and assayed on day 10 or later. The plates were subcultured and maintained in T cell growth medium

and stimulated with γ -irradiated EW-Ag-876 LCLs. Initial assays were against type A and type B LCLs and subsequently against peptide-coated PHA-blasts or LCLs. Precursor frequencies were determined by a modification of the method of maximum likelihood estimation (11).

- 5 Assessment of peptide specific activity. For determination of the antigenic specificity at the peptide level, a set of 20-mer overlapping peptides of EBNA-6B (residues 100-1069 (46)) was prepared using a kit and software distributed by Chiron Mimotopes (Chiron Corporation, Sydney, Australia). Nine-mer and 10-mer overlapping peptides with unblocked C and N termini
10 were manufactured by Chiron Mimotopes. Labelled target cells were preincubated with overlapping peptides in the 96-well trays for 30-60 minutes before adding effectors. Alternatively, labelled targets were preincubated with a high dose of peptide (40-80 μ g/ml) then washed twice before using in CTL assays. The washing procedure removes the
15 complication of bystander killing in the latter type of experiments (6).

Depletion experiments

- T-cell cultures were depleted of either CD4 or CD8 cells by pretreating with saturating levels of OKT4 (mouse anti-CD4) or OKT8 (mouse anti-CD8) for 30 min on ice before washing twice and incubating
20 with washed Dynabeads M450 (Sheep anti-mouse IgG coated beads, Dynal, A.S, Oslo, Norway) at 4°C for 60 min with rotation. Beads and adherent cells were removed magnetically and the remaining cells washed once before use in a CTL assay or FACScan analysis.

RESULTS

25

EBV typing of lymphocyte donor

- PCR analysis revealed the presence of both type A and type B EBV in blood from donor EW. The presence of both virus types was supported by the observation that both type A and type B spontaneous cell lines grew out
30 although only the former (referred to as EW-spon) was established in long term culture.

Selection of T-cell cultures for analysis

- Regression analysis indicated that the donor had memory CTL activity to both virus types. The number of cultures that could be assessed
35 was therefore minimal compared with the T cell responses of most type A donors investigated by this laboratory. Accordingly, 25 colonies from the

third sample were assayed and seven of these selected for further study. Three of these cultures (B4, B5 and B7) were type B specific i.e. they lysed autologous type B cells (>15% lysis) but not type A (<10 % lysis); the other four (A3, A9, B6, and B9) were A-B specific i.e. they lysed both type A and type B autologous LCLs (>15%) (Table 1).

CTL analysis of vaccinia-infected targets

The seven selected cultures were assayed for CTL activity towards vaccinia-infected autologous type A LCLs (EW-Wil). Four cultures (A3, A9, B4 and B5) gave significant enhancement of CTL activity when vacc-EBNA 6B infected LCLs were compared with control cells or vacc-TK infected target cells. Furthermore, these cultures reacted poorly with other EBV sequences (EBNA-1, EBNA-2B, EBNA-3B, EBNA-4A, LMP1, LMP2, BZLF1 and BHRF1) expressed by recombinant vaccinia viruses thus indicating specificity for the vacc-EBNA-6B infected target. Cultures B5 (type B specific) and A3 (crossreactive) were selected for more detailed analysis.

The specificity of cultures A3 and B5 was confirmed in a separate experiment in which EW-Wil target cells were infected with vacc-EBNA-6A and vacc-EBNA-6B. While the CTL activity of colony A3 was enhanced by infection of the target cells with either construct (<10% to 20-37%) the effector function of colony B5 was enhanced only when target cells were infected with vacc-EBNA-6B (<10% to 73-79.5%) (Fig. 2). These results suggest that culture B5 recognised a type B specific epitope within EBNA-6 while culture A3 recognised a crossreactive epitope within EBNA-6.

Identification of specific peptide targets of CTL activity

The type B specific culture B5 recognised the peptide corresponding to EBNA-6B residues 210-229 (peptide 12 of the EBNA-6B set) when assayed against individual 20-mer overlapping peptides preincubated with autologous PHA blasts (Fig. 3a). Nine-mer and 10-mer overlapping peptides were then assayed to identify the minimal epitope as the 10-mer QNGALAINTF (residues 213-222; Fig. 3b). The corresponding peptide from the type A overlapping sequence was not recognised by this culture, thus confirming the type B specificity of these T cells.

Culture A3 was assayed against pools of overlapping 20-mer peptides from EBNA 6B on autologous PHA-blasts in order to identify its peptide target. A pool containing 6 x 20-mer peptides (peptides 19-24 in the EBNA-6B set) from residue 280-349 was recognised by this CTL culture and these

peptides were subsequently assayed individually to identify the reactive 20-mer as peptide 19, residues 280-299 (Fig. 4a). In Fig. 4b, the minimal epitope was identified as LLDFVRFMGV corresponding to residues 284 -293 of the EBNA 6A and 6B sequences. This sequence is common to both type A and B of EBV and overlaps with the sequence of the B44-restricted epitope EENLLDFVRF (residues 281 - 290 of EBNA6A and 6B: (7)).

HLA-restriction of epitopes.

(a) Type B specific epitope

As Facs assays revealed that cultures recognising the EBNA 6B-encoded sequence QNGALAINTF contained mixtures of CD4⁺, CD4⁺ CD8⁺ and CD8⁺ cells when assayed initially, the possibility that the CTL activity was actually due to a CD4 component needed to be eliminated. Several of these cultures also contained components that killed type A autologous targets, presumably because they were derived from the limiting dilution format and contained a mixture of cells. Depletion and inhibition experiments confirmed that the CD8⁺ cells were the active CTLs.

In the depletion experiments, Dynabeads were used to deplete either CD4⁺ or CD8⁺ cells from reactive cultures. When samples of three different epitope-reactive cultures were tested, depletion of the CD4⁺ cells (which would have included the CD4⁺ CD8⁺ component) did not reduce the CTL activity, but depletion of the CD8⁺ cells completely abrogated the epitope specific activity (Fig. 5). Facs analysis in one of these experiments confirmed that the active fraction contained the CD8⁺ cells and that both the CD4⁺ and the CD4⁺-CD8⁺ cells had been removed. Fig. 6 illustrates the inhibition of the specific CTL activity of the mixed cultures by anti-class I monoclonal antiserum but not by the anti-class II serum that was positive for DR, DP and DQ alleles. It was concluded therefore that this epitope was Class I-restricted.

In order to identify the restricting allele. LCLs, Burkitt's lymphoma cell lines or PHA blasts of known HLA specificity were incubated with minimalised peptides and washed before using as target cells in CTL assays. As Table 2 indicates, the only target cells killed by the QNGALAINTF specific CTLs were those that were both A201 and B62 (B15) positive. In contrast, targets that were A201 but not B62, and target cells that only shared A24 or B51 alleles with the donor were not killed. It was therefore

concluded that the epitope is B62 (B15)-restricted. B62 which comprises at least 5 subtypes has recently been reclassified as part of the B15 family.

(b) Cross-reactive epitope

5 In similar experiments, culture A3 consistently recognised cells bearing HLA-A201 but not cells bearing A24, B51, or B62. Furthermore, the CTL activity of this culture was inhibited by anti-class I antibodies but not by anti-class II antibodies. Finally, the A3 minimal epitope corresponds to the motif already identified for HLA-A201 restriction (39). It was therefore
10 concluded that the epitope LLDFVRFMGV corresponding to residues 284-293 in both EBNA-6A and EBNA-6B was HLA Class I A201 restricted.

Predominance of type B CTL response in limiting dilution cultures and bulk cultures

15 Limiting dilution culture methodology was used in order to investigate the magnitude of the type B specific CTL response. PBL stimulated by a feeder layer of irradiated autologous type B cells were assayed after 10 days against type A and type B autologous LCLs. The results indicated a higher frequency of T cell precursors killing type B LCLs
20 than of those killing type A LCLs. There were therefore a number of wells at limiting dilutions that were type B specific. In general, the wells manifesting type B specific activity retained this specificity. After minimalising the epitopes as described above, the cultures were retested for epitope specific activity. A frequency of 1/16395 was calculated for the Type
25 B epitope QNGALAINTF compared with 1/304,890 for the A-B specific epitope LLDFVRFMGV when these cultures were assayed 12 weeks later.

Additional EBV CTL epitopes were obtained from other donors using similar methods.

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DISCUSSION

The identification of an EBV type B specific epitope represents the first demonstration of a CTL epitope encoded within type B specific regions
35 of the EBV genome. The epitope was identified as QNGALAINTF corresponding to residues 213-222 of EBNA-6B. The collective data

presented here indicates that stimulation of T cells from donor EW by autologous type B infected LCLs reactivated a predominantly type B response rather than a cross-reactive response. The specificity of the type B response could be maintained on extended culture *in vitro*. Clearly the response to QNGALAINTF would seem to be an immunodominant one as indicated by the number of reactive wells in the limiting dilution cultures. It is also interesting to note that a number of type B- specific CTL cultures were established for which specific epitopes could not be identified. It is possible that the target epitopes for these T cells lie within EBNA-4B or outside the regions included in the EBNA-2B, -3B and -6B constructs.

The epitope QNGALAINTF appears to be restricted through HLA B62 (B15), although the subtype specificity has yet to be determined. Motifs for B*1501 have previously been published (39) and F is commonly found in the last anchor position (9 or 10) as in the epitope identified here. Q or L appear to be the preferred residues at position 2 of B*1501; for the new epitope, Q is at position 1. The corresponding type A sequence for this peptide (QNAARTLNTF) was nonreactive in CTL assays, thus confirming the type B specificity of the epitope.

A second epitope LLDFVRFMGV also represents a previously unrecognised EBV-encoded epitope. This sequence corresponds to residues 284 -293 of both EBNA-6A and 6B and was shown to be restricted by HLA-A201.

An interesting facet of this work was the detection of both type A and type B EBV in the one donor by PCR of both PBL and spontaneous cell lines. Although the available evidence indicates a high prevalence of type B in the community, it is known that in healthy donors type A predominates as the one that is easiest to isolate. Until now, responses to type A virus have therefore been easier to identify.

It was noted in the results section that the donor did not give a particularly active cloning response. It is possible that the type B viruses are less immunogenic than type A, or even that type B EBV has evolved in part to evade the immune system. Type B viruses could conceivably normally be resident in the epithelial compartment rather than the lymphocyte compartment due to their lower transformation efficiency. In this situation, type B epitopes may not be readily presented to the immune system. As already implied, the lower transformation efficiency of type B viruses (42)

may hinder reinforcement over time of the immune response to the latency antigens.

Finally, it is important when considering the possibility of vaccinating against EBV to recognise that it may be necessary to vaccinate against both type A and type B. If a patient, particularly a transplant patient at risk of EBV-related post-transplant lymphoma (8), is protected against type A but not type B, there may even be greater risk of clinical disease if subsequently infected with the other virus type. It is particularly important to consider EBV type B epitopes in view of the finding that type B is commonly detected in immunosuppressed individuals. Subunit vaccines against EBV are currently being trialed. In the long term, such vaccines would ideally contain cross-protective specificities. Definition of type B and cross-reactive epitopes allows development of a more effective vaccine, more effective in terms of the range of EBV specificities and HLA types that are covered by the vaccine.

It will be appreciated by persons skilled in the art that numerous variations and/or modifications may be made to the invention as shown in the specific embodiments without departing from the spirit or scope of the invention as broadly described. The present embodiments are, therefore, to be considered in all respects as illustrative and not restrictive.

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Table 1. Initial Selection of CTL Cultures

Culture No. #	Target Cells (% Lysis)		K562 (NK sensitive)		Facs Markers (% + ve)				Specificity *
	FW-Wil (type A)	FW-Ag-876 (type B)	876	8.2	CD3 ⁺	CD4 ⁺	CD8 ⁺	CD4 ⁺ CD8 ⁺	
A3	33.8	17.0		8.2	97.5	0.3	85	2	A-B
A9	17.7	10.0		6.1	89	2	82	10	A-B
B4	6.6	16.6		4.3	93	2	70	0	B
B5	5.9	23.8		7.8	95	4	90.5	5.1	B
B6	18.5	31.3		9.6	80	2	66	7	A-B
B7	8.1	30.2		4.1	86	80	0	1	B
B9	16.8	7.7		4.1	87	0	96	0	A-B

Cultures designated A were generated by stimulation with type A autologous LCLs; cultures designated B were derived by stimulation with type B autologous LCLs

* A-B specificity: > 10% of both type A and B autologous cell lines; type B specificity: < 10% lysis of type A and > 15% lysis of type B autologous cell lines

Table 2. HLA-Restriction of type B-specific and crossreactive epitopes

TARGET CELLS (Transforming isolate)	EBV TYP E	CLASS I ALLELES	B-type epitope CTL		Cross-reactive epitope CTL	
			-Pep	+Pep	-Pep ²	+Pep
EW-PHA BLASTS	-	A2 A24 B51 B62	-	+	-	++
RM-LCL (WIL)	A	A101 A2 B8 B62	-	+	-	++
LP-LCL (WIL)	A	A2 A32 B35 B62	-	+	+-	++
PGP-LCL (PUY)	A	A1 A24 B8 B14	-	-	-	-
WW II-LCL (WWII)	A	A11 A24 B18 B39	-	-	-	((+))
IM-LCL (GOR)	B	A1 A11 B51 B8	-	NT	-	NT
AS-LCL	A,B	A201 A24 B51 B62	-	NT	NT	NT
SB-PHA	-	A201 A201 B35 B57	-	-	-	++
CHEP (BL) ¹	A	A2 A3 B7	-	-	-	-
MUTU (BL)	A	A1 A2 B45	-	-	-	+
TM-LCL (Wil)	A	A11 A32 B35 B15	-	-	-	-
IAP-LCL (Wil)	A	A2 A28 B8 B15	-	+	-	+

1: BL = Burkitt's Lymphoma cell lines - used as targets in peptide experiments as the level of background killing is low

2: Clone A3 was originally selected on the basis of CTL activity towards both EW-Wil (type A) and EW-Ag-876 (type B) LCL's, but in the above experiments this clone had low activity only towards heterologous Wil-transformed LCL's.

Cultures that recognised either the crossreactive epitope or the type B-specific epitope were tested in CTL assays against heterologous cell lines with or without specific peptide

Table 3

EBV Antigen	Peptide Epitope	HLA Restriction	EBV Type
EBNA3A	QVKWRMTTL	HLA B8	Type 2
EBNA3A	VFSDGRVAC	HLA A29	Type 1
EBNA3A	VPAPAGPIV	HLA B7	Type 1
EBNA3B	TYSAGIVQI	HLA A24	Type 1
EBNA3C	LLDFVRFMGV	HLA A2	Type 1 & 2
EBNA3C	QNGALAINTF	HLA B62	Type 2
EBNA3A	VSSDGRVAC	HLA A29	Type 1
EBNA3A	VSSEGRVAC	HLA A29	Type 1
EBNA3A	VSSDGRVPC	HLA A29	Type 1
EBNA3A	VSSDGLVAC	HLA A29	Type 1
EBNA3A	VSSDGQVAC	HLA A29	Type 1
EBNA3A	VSSDGRVVC	HLA A29	Type 1
EBNA3A	VPAPPVGPIV	HLA B7	Type 1
EBNA3A	VEITPYEPTG	HLA B44	Type 1
EBNA3A	VEITPYEPTW	HLA B44	Type 1
EBNA3A	VELTPYKPTW	HLA B44	Type 1
EBNA3C	RRIYDLIKL	HLA B27	Type 1
EBNA3C	RKIYDLIEL	HLA B27	Type 1
LMP2A	PYLFWLAGI	HLA A23	

Table 4

EBV Antigen	Peptide Epitope	HLA Restriction	EBV Type	Reference
EBNA1	TSLYNLRRGTALA	HLA DR1	Type 1 & 2	(Khanna 1995A)
EBNA2	DTPLIPLTIF	HLA B51/A2	Type 1	(Schmidt 1991)
EBNA2	TVFYNIIPPMPL	HLA DQ2	Type 1	WO 95/24925
EBNA3A	VEITPYKPTW	HLA B44	Type 1	WO 95/24925
EBNA4	VSFIEFVGW	HLA B57	Type 1	WO 95/24925
EBNA6	FRKAQIQGL	HLA B57	Type 1	WO 95/24925
EBNA3A	FLRGRAYGL	HLA B8	Type 1	(Burrows 1992)
EBNA3A	QAKWRLQTL	HLA B8	Type 1	(Burrows 1994B)
EBNA3A	SVRDRLARL	HLA A2	Type 1 & 2	(Burrows 1994B)
EBNA3A	YPLHEQHGM	HLA B35	Type 1	(Burrows 1994B)
EBNA3A	HLAAQGMAY	HLA?	Type 1	(Burrows 1994B)
EBNA3A	RPPIFIRRL	HLA B7	Type 1	(Hill 1995)
EBNA3A	RLRAEAGVK	HLA A3	Type 1	(Hill 1995)
EBNA3B	IVTDFSVIK	HLA A11	Type 1	(Gavioli 1993A)
EBNA3B	AVFDRKSDAK	HLA A11	Type 1	(Gavioli 1993A)
EBNA3A	*NPTQAPVIQLVHAVY	HLA A11	Type 1	(Gavioli 1993A)
EBNA3A	*LPGPQVTAVLLIIEES	HLA A11	Type 1	(Gavioli 1993A)
EBNA3A	*DEPASTEPVHDQIL	HLA A11	Type 1	(Gavioli 1993A)
EBNA3A	RYSIFFDY	HLA 24	Type 1	(Burrows 1992)
EBNA3B	AVLLHEESM	HLA B35	Type 1	(Khanna 1995)
EBNA3B	RRARSLAERY	HLA B27	Type 1	(Hill 1995)
EBNA3C	EENLLDFVRF	HLA B44	Type 1 & 2	(Burrows 1990)
EBNA3C	KEHVIQNAF	HLA B44	Type 1	(Khanna 1992)
EBNA3C	RRYDLIEL	HLA B27	Type 1	(Brooks 1993B)
EBNA3C	QPRAPIRPI	HLA B7	Type 1 & 2	(Hill 1995)
EBNA3C	EGGVGWRHW	B44		(Morgan 1996)
LMP2A	CLGGLTMTV	HLA A2	Type 1 & 2	(Lee 1993B)
LMP2A	RRRWRLTV	HLA B27	Type 1	(Brooks 1993B)
Zta	RAKFKQLL	HLA B8		(Bogedain 1995)
Zta	*RKCCRAKFKQLLQHYR	HLA Cw6		(Bogedain 1995)

References and legend for Table 4

* Epitope sequence not minimalised

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- 35

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CLAIMS:

1. A cytotoxic Epstein-Barr virus T-cell epitope, the epitope being selected from the group consisting of QVKWRMTTL, VFSDGRVAC, VPAPAGPIV, TYSAGIVQI, LLDFVRFMGV, QNGALAINTF, VSSDGRVAC, VSSEGRVAC, VSSDGRVPC, VSSDGLVAC, VSSDGQVAC, VSSDGRVVC, VPAPPVGPIV, VEITPYEPTG, VEITPYEPTW, VELTPYKPTW, RRIYDLIKL, RKIYDLIEL and PYLFWLAGI.
2. A subunit vaccine, the vaccine including at least one T-cell epitope selected from the group consisting of QVKWRMTTL, VFSDGRVAC, VPAPAGPIV, TYSAGIVQI, LLDFVRFMGV, QNGALAINTF, VSSDGRVAC, VSSEGRVAC, VSSDGRVPC, VSSDGLVAC, VSSDGQVAC, VSSDGRVVC, VPAPPVGPIV, VEITPYEPTG, VEITPYEPTW, VELTPYKPTW, RRIYDLIKL, RKIYDLIEL and PYLFWLAGI.
3. A vaccine as claimed in claim 2 in which the subunit vaccine includes at least one further epitope selected from the group consisting of TSLYNLRRGTALA, DTPLIPLTIF, TVFYNNIPMPL, VEITPYKPTW, VSFIEFVGW, FRKAQIQGL, FLRGRAYGL, QAKWRLQTL, SVRDRLARL, YPLHEQHGM, HLAAQGMAY, RPPIFIRRL, RLRAEAGVK, IVTDFSVIK, AVFDRKSDAK, NPTQAPVIQLVHAVY, LPGPQVTAVLLHEES, DEPASTEPVHDQLL, RYSIFFDY, AVLLHEESM, RRARSLSAERY, EENLLDFVRF, KEHVIQNAF, RRIYDLIEL, QPRAPIRPI, EGGVGWRHW, CLGGLTMMV, RRRWRRLTV, RAKFKQLL and RKCCRAKFKQLLQHRY.
4. A vaccine as claimed in claim 2 or claim 3 in which the subunit vaccine includes the cytotoxic T-cell epitopes LLDFVRFMGV, QVKWRMTTL and FLRGRAYGL.
5. A vaccine as claimed in claim 2 or claim 3 in which the subunit vaccine includes the cytotoxic T-cell epitopes LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL and EENLLDFVRF.
6. A vaccine as claimed in claim 2 or claim 3 in which the subunit vaccine includes the cytotoxic T-cell epitopes LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL and QPRAPIRPI.
7. A vaccine as claimed in claim 2 or claim 3 in which the subunit vaccine includes the cytotoxic T-cell epitopes LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL, EENLLDFVRF and QPRAPIRPI.
8. A vaccine as claimed in any one of claims 2 to 7 in which the vaccine further includes at least one antigen to which the individual will

mount an anamnestic response in addition to the at least one cytotoxic T-cell epitope.

9. A vaccine as claimed in claim 8 in which the at least one antigen is selected from the group consisting of tetanus toxoid, diphtheria toxoid,
5 Bordetella pertussis antigens, poliovirus antigens, purified protein derivative (PPD), gp350 protein, helper epitopes and combinations thereof.
10. A vaccine as claimed in claim 9 in which the at least one antigen is tetanus toxoid.
11. A vaccine as claimed in any one of claims 2 to 10 in which the
10 vaccine comprises a water-in-oil formulation
12. A nucleic acid vaccine, the vaccine including a nucleic acid sequence encoding at least one of the cytotoxic T-cell epitopes selected from the group consisting of QVKWRMTTL, VFSDGRVAC, VPAPAGPIV, TYSAGIVQI, LLDFVRFMGV, QNGALAINTF, VSSDGRVAC, VSSEGRVAC,
15 VSSDGRVPC, VSSDGLVAC, VSSDGQVAC, VSSDGRVVC, VPAPPVGPIV, VEITPYEPTG, VEITPYEPTW, VELTPYKPTW, RRIYDLIKL, RKIYDLIEL and PYLFWLAGI.
13. A nucleic acid vaccine as claimed in claim 12 in which the nucleic acid sequence encodes at least one additional cytotoxic T-cell epitope
20 selected from the group consisting of TSLYNLRRGTALA, DTPLIPLTIF, TVFYNIPPMPL, VEITPYKPTW, VSFIEFVGW, FRKAQIQGL, FLRGRAYGL, QAKWRLQTL, SVRDLARL, YPLHEQHGM, HLAAQGMAY, RPIFIRRL, RLRAEAGVK, IVTDFSVIK, AVFDRKSDAK, NPTQAPVIQLVHAVY, LPGPQVTAVLLHEES, DEPASTEPVHDQLL, RYSIFFDY, AVLLHEESM,
25 RRARSLSAERY, EENLLDFVRF, KEHVIQNAF, RRIYDLIEL, QPRAPIRPI, EGGVGWRHW, CLGGLTMV, RRRWRRLTV, RAKFKQLL and RKCCRAKFKQLQHYP.
14. A nucleic acid vaccine as claimed in claim 12 or claim 13 in which the nucleic acid sequence encodes LLDFVRFMGV, QVKWRMTTL and
30 FLRGRAYGL, or LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL and EENLLDFVRF, or LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL and QPRAPIRPI, or LLDFVRFMGV, QVKWRMTTL, FLRGRAYGL, EENLLDFVRF and QPRAPIRPI.

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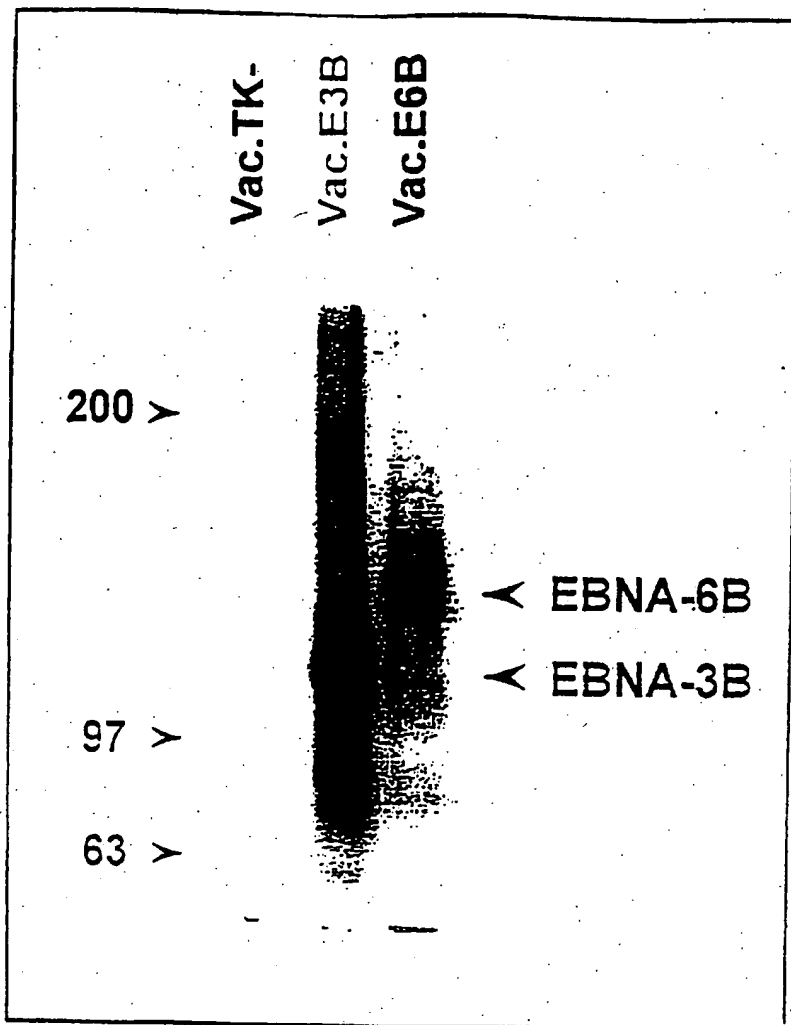


FIGURE 1

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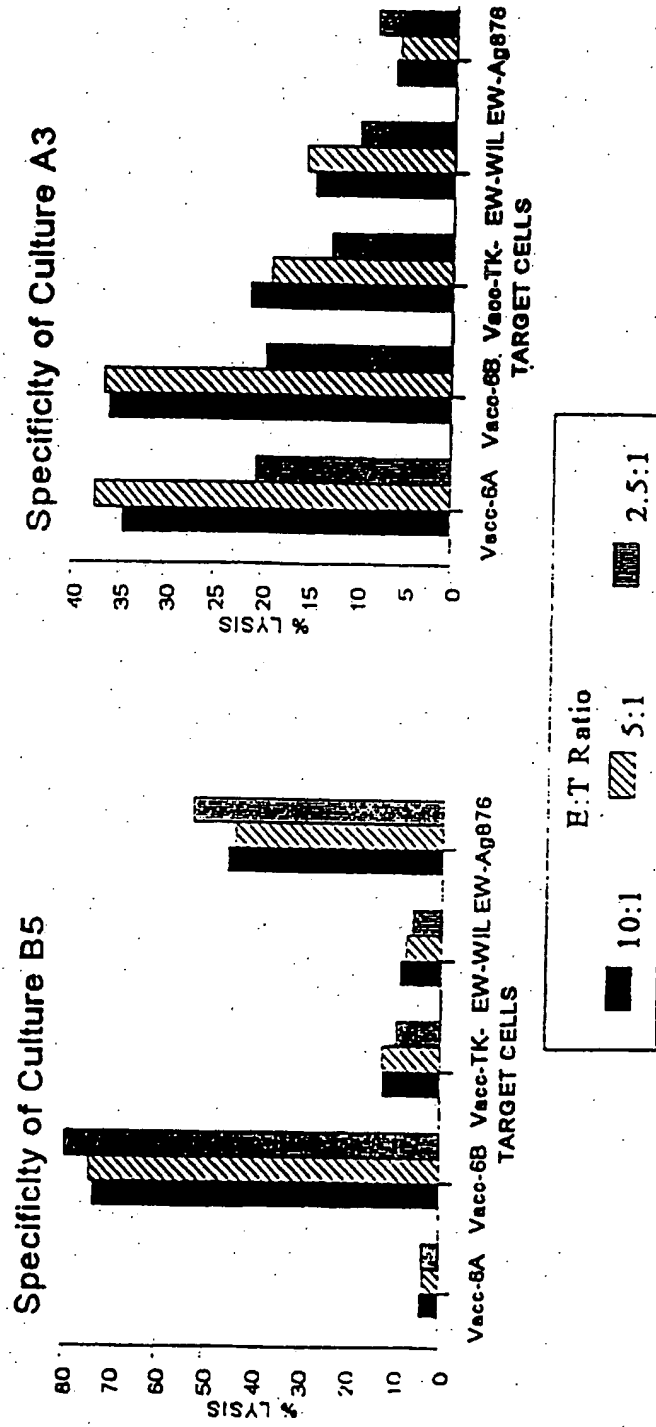


FIGURE 2

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Fig. 3a

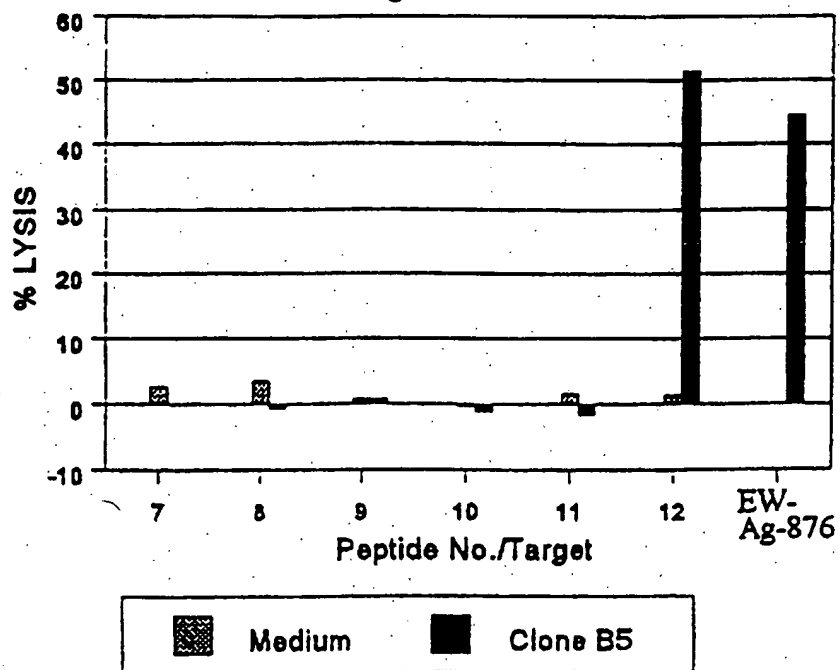
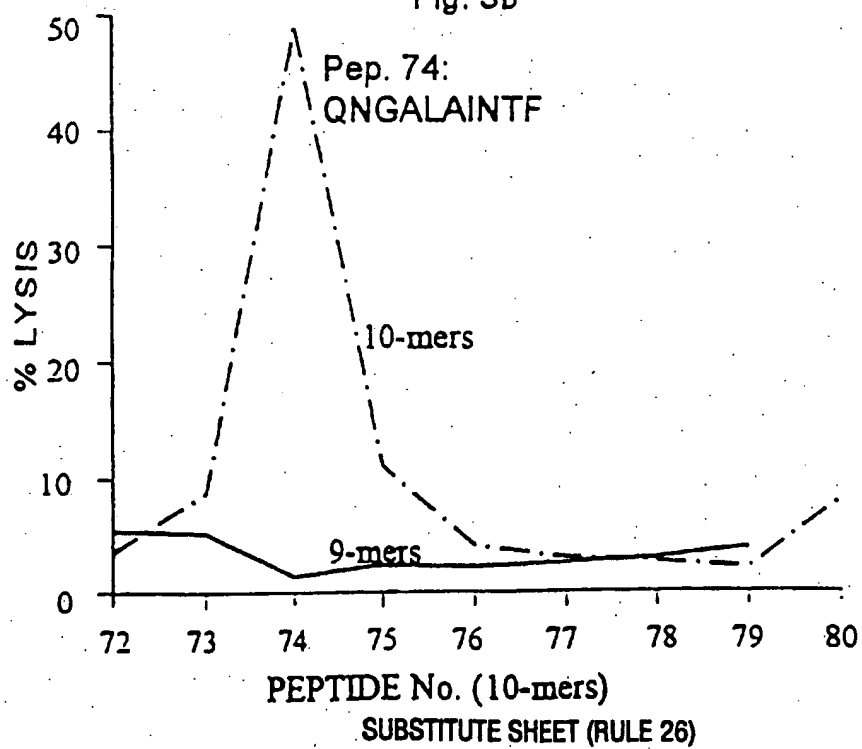


Fig. 3b



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Fig. 4a

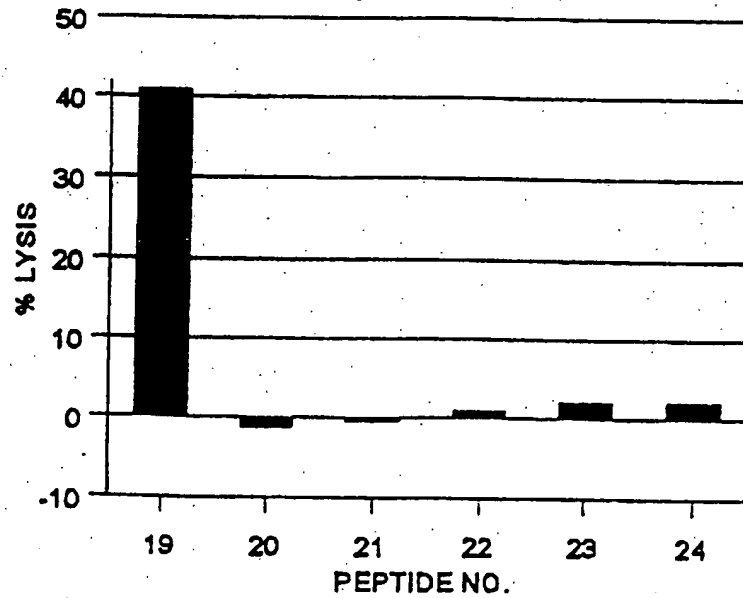
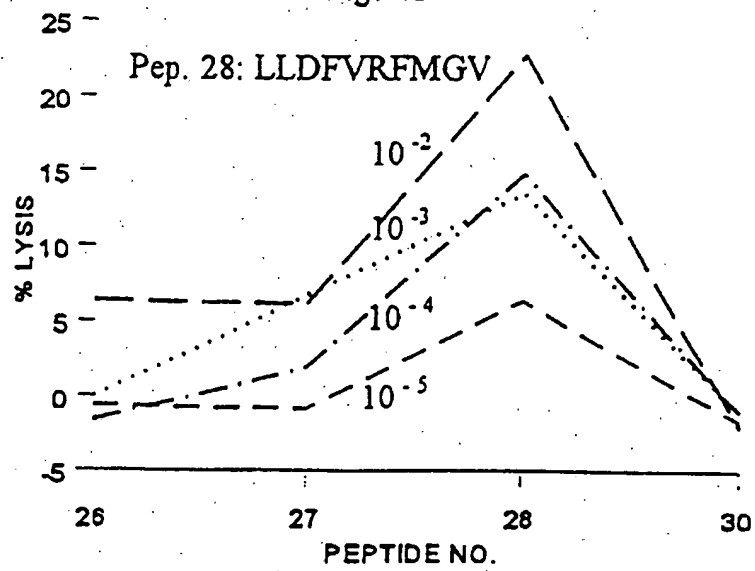
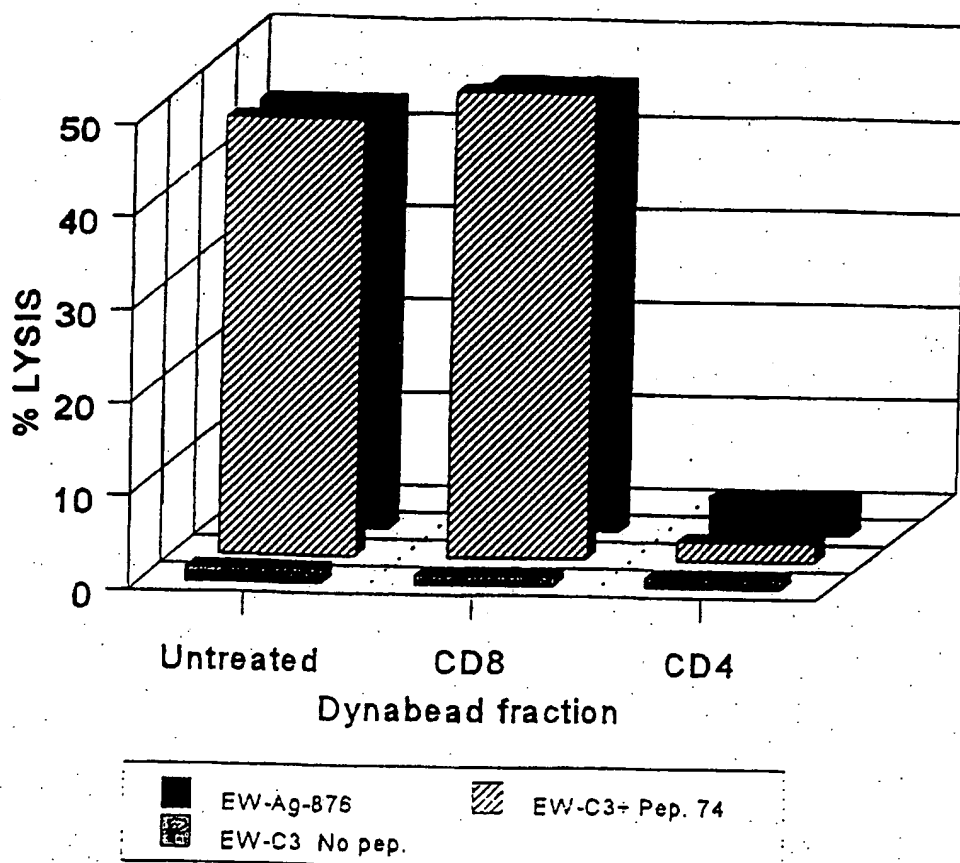


Fig. 4b



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Figure 5

SUBSTITUTE SHEET (RULE 26)

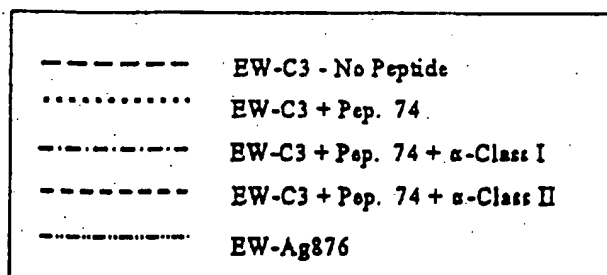
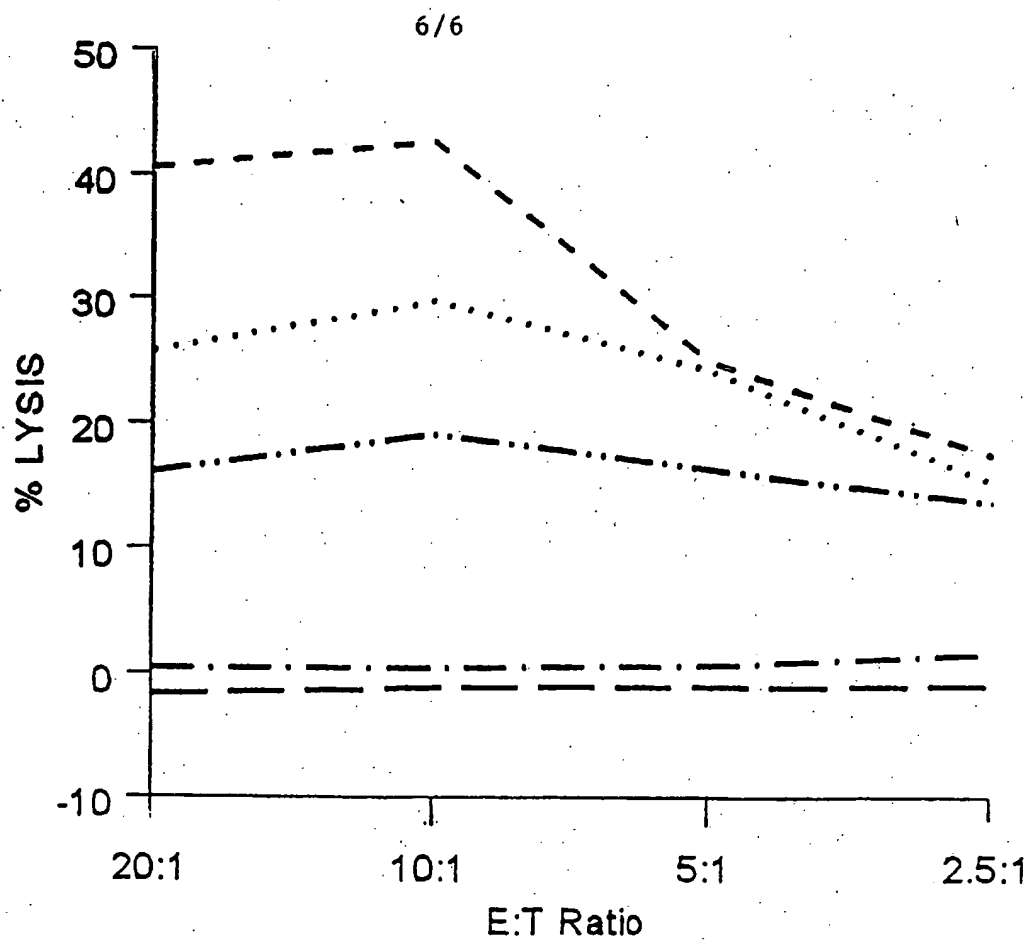
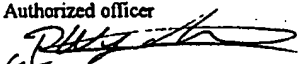


FIGURE 6

INTERNATIONAL SEARCH REPORT

International Application No.
PCT/AU 97/00328

A. CLASSIFICATION OF SUBJECT MATTER												
Int Cl ⁶ : C07K 007/06 A61K 039/245												
According to International Patent Classification (IPC) or to both national classification and IPC												
B. FIELDS SEARCHED												
Minimum documentation searched (classification system followed by classification symbols) See Electronic Database Box below												
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched See Electronic Database Box below												
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) Sequence search in Chem Abs and DGENE as follows qkvwrmttl vbsdgrvac vpapagpiv tysagivqi lldfvrimgv qngalaintf vssdgrvac vssegrvac vssdgrvpc vssdglvac vssdgvvac vssdgrvvc vpappvgpiv veitpyeptg veitpyeptw veltpykptw rriydlkl rkiydlkl pylfwlagi.												
C. DOCUMENTS CONSIDERED TO BE RELEVANT												
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.										
P,X	<u>Journal of Virology</u> , December 1996, volume 70, number 12, pages 8858-8864. KERR, B.M., (et al) "Identification of Type B-specific and Cross-Reactive Cytotoxic T-Lymphocyte Responses to Epstein-Barr Virus".	1-14										
A	<u>Journal of Virology</u> , August 1990, volume 64, number 8, pages 3974-3976. BURROWS, S.R. (et al) "An Epstein-Barr Virus-Specific Cytotoxic T-Cell Epitope Present on A- and B-type Transformants".											
<input type="checkbox"/> Further documents are listed in the continuation of Box C <input type="checkbox"/> See patent family annex												
<p>* Special categories of cited documents:</p> <table border="0"> <tr> <td>"A" document defining the general state of the art which is not considered to be of particular relevance</td> <td>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</td> </tr> <tr> <td>"E" earlier document but published on or after the international filing date</td> <td>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</td> </tr> <tr> <td>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</td> <td>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</td> </tr> <tr> <td>"O" document referring to an oral disclosure, use, exhibition or other means</td> <td>"&" document member of the same patent family</td> </tr> <tr> <td>"P" document published prior to the international filing date but later than the priority date claimed</td> <td></td> </tr> </table>			"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention	"E" earlier document but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone	"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art	"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family	"P" document published prior to the international filing date but later than the priority date claimed	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention											
"E" earlier document but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone											
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art											
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family											
"P" document published prior to the international filing date but later than the priority date claimed												
Date of the actual completion of the international search 20 June 1997		Date of mailing of the international search report 25 JUN 1997										
Name and mailing address of the ISA/AU AUSTRALIAN INDUSTRIAL PROPERTY ORGANISATION PO BOX 200 WODEN ACT 2606 AUSTRALIA Facsimile No.: (06) 285 3929		Authorized officer  J.H. CHAN Telephone No.: (06) 283 2340										